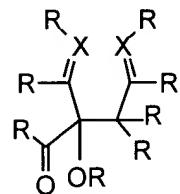
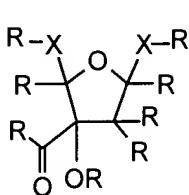


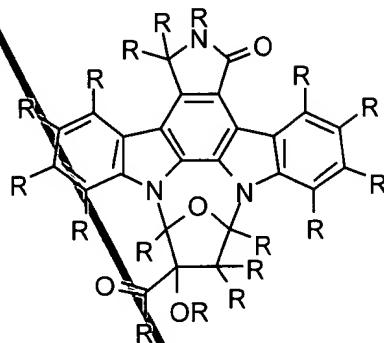
C1
cont

with an acetal selected from the group consisting of the formulae



A2

and mixtures thereof,
to produce a glycosylated product of the formula

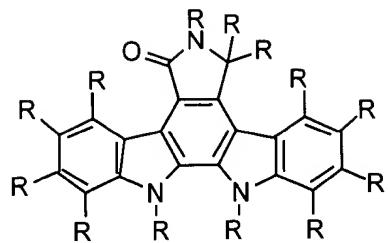


wherein R is selected from the group consisting of

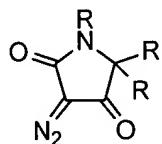
- a) a C₃₋₁₀ branched or unbranched alkyl, optionally partially or fully halogenated, hydroxy, C₁₋₃ alkyloxy, carboxy, amino, alkylamino, including Me, CH₂OH, and CO₂ Me;
 - b) an aryl optionally substituted with one to five groups consisting of halo, hydroxy, C₁₋₃ alkyloxy, including Bn, DMB, and PMB;
 - c) a hydrogen;
 - d) a halogen; and
 - e) mixtures of any of these, and
- wherein X is S and/or O.

AB
C5-

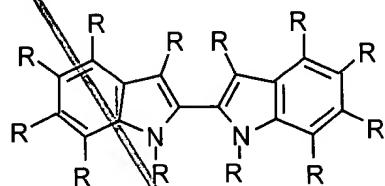
17. A process for the preparation of furanosylated indolocarbazoles by first preparing an indolocarbazole of the formula



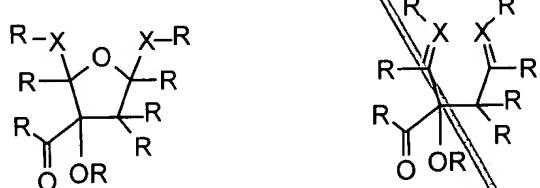
by reacting a diazo compound of the formula



with a biindole of the formula



*C5
ABX
w*
in the presence of a transition metal catalyst in a solvent capable of solvating the reactants, and then reacting the indolocarbazole with an acetal selected from the group consisting of the formulae



and mixtures thereof, in the presence of a Bronsted acid or a Lewis acid to produce a glycosylated product of the formula